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(71) Applicant: BIO-MEGA INC.
2100 Rue Cunard
Laval Québec H7S 2G5(CA)

(72) Inventor: Anderson, Paul Cates
4130 Edward Higgins
Pierrefonds, Quebec, H8Y 3M8(CA)
Inventor: Guindon, Yvan
12230 Le Mesurier, Montreal
Quebec, H4K 2B3(CA)
Inventor: Yoakim, Christiane
63 Gianchetti
Laval, Quebec, H7G 4T5(CA)

(74) Representative: Laudien, Dieter, Dr. et al
Boehringer Ingelheim GmbH, Abteilung
Patente
D-6507 Ingelheim am Rhein(DE)

(54) Enzyme inhibitors.

(57) Disclosed herein are peptide derivatives which inhibit the activity of human immunodeficiency virus (HIV) protease. The peptide derivatives can be represented by general formula $R^1-R^2-Y-R^3-R^4$ wherein R^1 is an optionally substituted 3-(1,2,3,4-tetrahydroisoquinolyl)carbonyl residue, R^2 and R^3 are amino acid or analogous amino acid residues (R^3 may optionally be absent), Y is a non-peptide linking unit, e.g. statyl, and R^4 is $[-NR^{17}CH(R^{18})C(O)]_p-Z$ wherein R^{17} is hydrogen or lower alkyl, R^{18} is an amino acid or analogous amino acid side chain, p is zero or 1 and Z is a terminal group (e.g. hydroxy or amino), or R^4 is $-NR^{17}CR^{18}(R^{21})CH_2OH$ wherein R^{17} and R^{18} are as noted hereinabove and R^{21} is hydrogen, lower alkyl or hydroxy(lower)alkyl. The derivatives also inhibit renin activity. Accordingly, the derivatives can be used for combating HIV infections or for treating hypertension or congestive heart failure.

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FIELD OF THE INVENTION

This invention relates to compounds having valuable pharmacological properties. More specifically, the invention relates to peptide derivatives (hereinafter called "peptides") exhibiting activity against particular retroviruses, to processes for producing the peptides, to pharmaceutical preparations thereof, and to a method of using the peptides to combat infections caused by the retroviruses. The invention also relates to pharmaceutical preparations and a method of treating renin-dependent hypertension and congestive heart failure wherein the peptides serve as the active agent.

BACKGROUND OF THE INVENTION

During the last ten years, retroviruses have emerged from relative obscurity to prominence. These viruses now are known to cause of variety of diseases in vertebrates, the most insidious to humans being immunodeficiencies and cancers.

In 1983, a retrovirus, known as human immunodeficiency virus type 1 (HIV-1), was established as a causative agent for acquired immune deficiency syndrome (AIDS). This virus has become a pestilence of alarming proportion. More recently, the closely related virus, human immunodeficiency virus type 2 (HIV-2) has been identified as a second causative agent of AIDS. Hereinafter, the term "HIV" is meant to include both HIV-1 and HIV-2 and any mutants thereof.)

Presently, several compounds are being evaluated in the clinic as possible therapeutic agents for AIDS. Another compound, 3-azido-3-deoxythymidine (known also as zidovudine or AZT), has been shown in the clinic to decrease mortality and the frequency of opportunistic infections in AIDS patients. This latter compound is being used to manage certain patients with symptomatic HIV infections. However, in spite of some recent progress, the need for an effective therapy for AIDS still exists. For recent reviews, see R.A. Weiss in "Molecular Basis of Virus Disease", Symposium of the Society for General Microbiology, Vol. 40, Eds. W.C. Russel and J.W. Almond, University Press, Cambridge, UK, 1987, pp 167-192, and R.C. Gallo and L. Montagnier, Scientific American, 259, (4), 40 (1988).

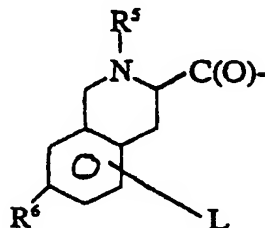
One approach to finding agents having anti-HIV activity is to inhibit the action of HIV-encoded enzymes. This manner of inhibition interferes with the replication and propagation of the virus. Such an approach has been applied successfully in a search for inhibitors of the viral encoded enzyme, reverse transcriptase (RT). More explicitly, the previously noted zidovudine was found to inhibit RT which is required to effect viral replication. Subsequently, zidovudine was developed as an anti-HIV agent. Still more recently, this approach has been investigated using another HIV-encoded enzyme known as HIVprotease as the target enzyme. In one instance, pepstatin A was found to inhibit the intracellular processing that provides the requisite HIV protease. See, S. Seelmeier et al., Proc. Natl. Acad. Sci. USA, 85, 6612 (1988). However, the development of pepstatin A as an anti-HIV agent seems improbable in view of its multiple activities. In another instance, M.L. Moore et al., Biochem. Biophys. Res. Comm., 159, 420 (1989), reported on investigations showing the inhibition of HIV protease by three heptapeptide analogs modeled after a conserved cleavage site (gag gene region) of the viral genomic polyprotein. A.D. Richards et al., FEBS Letters 247, 113(1989), also have reported that acetylpepstatin and nonapeptide analog inhibit HIV protease in vitro.

The present application discloses a group of peptide derivatives which are potent inhibitors of HIV protease and renin. These attributes, together with the attributes of a relatively selective action and an apparent lack of toxicity, renders the peptides useful as agents for combating HIV infections and for treating renin-associated hypertension and congestive heart failure.

The present peptides of this application are distinguished readily from pepstatin A and the previously noted peptide analogs by chemical and biochemical differences. The present peptides also possess a partial structural resemblance to peptide derivatives reported to be renin inhibitors; for instance, see D.F. Veber et al., European patent application 77,028, published April 20, 1983, and A. Wagner et al., Australian patent application 76241/87, published February 4, 1988. The remaining structural features and differences in biological profiles distinguish these latter prior art compounds from the present peptide derivatives, notwithstanding the existence of broad generic disclosures, such as R. Ten Brink, PCT patent application WO87/02986, published May 21, 1987, encompassing a myriad of compounds ranging in the millions. Finally, a class of peptide isosteres have been reported recently to have the unusual combination of renin inhibitory and antiretroviral activities; the latter compounds have structures which are quite different from the present peptides (see B. Weldmann, UK patent application 2203740, published October 26, 1988).

SUMMARY OF THE INVENTION

The peptides of this invention are represented by formula 1 $R^1-R^2-Y-R^3-R^4$ 1
 wherein R^1 is a derived amino acid radical of formula 2



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wherein R^5 is hydrogen, lower alkyl, or $QOC(O)-$ or $QC(O)-$ wherein Q is lower alkyl, lower cycloalkyl, (lower cycloalkyl)methyl, phenyl, phenyl monosubstituted with lower alkyl, lower alkoxy or halo, benzyl or benzyl monosubstituted with lower alkyl, lower alkoxy or halo; R^6 is hydrogen, hydroxy, lower alkyl, lower alkoxy, amino, nitro or $-OCH_2C(O)OR^7$ wherein R^7 is hydrogen, lower alkyl, lower cycloalkyl, (lower cycloalkyl)methyl, phenyl, phenyl monosubstituted with lower alkyl, lower alkoxy or halo, benzyl, benzyl monosubstituted with lower alkyl, lower alkoxy or halo; and L on the aromatic ring of the radical of formula 2 represents hydrogen or a substituent on the aromatic ring, the substituent being selected from the group of lower alkyl, lower alkoxy and halo, or L represents the same or different of two substituents on the aromatic ring, the substituents being selected from the group of lower alkyl, lower alkoxy and halo, provided that any two substituents do not interfere with each others presence;

R^2 is $-N(R^8)CH(R^9)C(O)-$ wherein R^8 is hydrogen or lower alkyl and R^9 is hydrogen, lower alkyl, lower alkyl monosubstituted with hydroxy, methoxy, methylthio or benzyloxy, lower cycloalkyl, (lower cycloalkyl)methyl, benzyl, 4-imidazolylmethyl, 2-thienylmethyl, 2-thiazolylmethyl, (4-hydroxyphenyl)methyl, [4-(lower alkoxy)phenyl]methyl, $-CH(OH)C_6H_5$, $-(CH_2)_4NH_2$, or $-(CH_2)_nC(O)OR^{1*}$ or $-(CH_2)_nC(O)NR^{11}R^{12}$ wherein n is the integer one, two or three, R^{1*} is hydrogen, a straight or branched chain alkyl containing one to ten carbon atoms, or phenyl(lower)alkyl and R^{11} , and R^{12} each independently is hydrogen, lower alkyl, phenyl-(lower)alkyl, or R^{11} , and R^{12} together with the nitrogen to which they are joined form a pyrrolidino, piperidino, morpholino, piperazino or 4-(lower alkyl)piperazino;

Y is a derived amino acid radical of the formula $-NHCH(R^{13})W-$ wherein R^{13} is lower alkyl, (lower cycloalkyl)methyl, $-CH_2CH_2SCH_3$, benzyl or benzyl substituted on the aromatic portion thereof with hydroxy or lower alkoxy, and W is $-CH(OH)-CH_2C(O)-$ or $-CH_2NCH(R^{14})C(O)-$ wherein R^{14} has the same meaning as defined for R^{13} ;

R^3 is absent or is $-N(R^{15})CH(R^{16})C(O)-$ wherein R^{15} is hydrogen or lower alkyl and R^{16} has the same meaning as defined herein for R^9 ; and

R^4 is $[-NR^{17}CH(R^{18})C(O)]_p-Z$ wherein R^{17} is hydrogen or lower alkyl, R^{18} has the same meaning as defined herein for R^9 ; p is the integer zero or one, and Z is hydroxy, lower alkoxy, benzyloxy or $-NR^{19}R^{2*}$ wherein R^{19} and R^{2*} each independently is hydrogen, lower alkyl or phenyl(lower)alkyl, or R^{19} and R^{2*} together with the nitrogen atom to which they are joined form a pyrrolidino, piperidino, morpholino, piperazino or 4-(lower alkyl)piperazino; or

R^4 is $-NR^{17}CR^{18}(R^{21})CH_2OH$ wherein R^{17} and R^{18} are as defined herein and R^{21} is hydrogen, lower alkyl or lower alkyl monosubstituted with hydroxy; or a therapeutically acceptable salt thereof.

A preferred group of the peptides of this invention for inhibiting HIV protease is represented by formula 1 wherein R^1 is the radical of formula 2 wherein R^5 is hydrogen, R^6 is hydrogen, hydroxy, methyl, methoxy, amino, nitro or $-OCH_2C(O)OR^7$ wherein R^7 is hydrogen, lower alkyl or benzyl, L is hydrogen or one or two halo substituents as defined herein; R^2 is $-N(R^8)CH(R^9)CO-$ wherein R^8 is hydrogen or methyl and R^9 is lower alkyl, lower cycloalkyl, cyclopropylmethyl, cyclohexylmethyl, $-CH_2CH_2COOH$, $-CH_2CH_2CONH_2$, $-CH_2CONH_2$ or benzyl; Y is $-NHCH(R^{13})W-$ wherein R^{13} is lower alkyl, (lower cycloalkyl)methyl, $-CH_2CH_2SCH_3$, benzyl or (4-methoxyphenyl)methyl and W is $-CH(OH)CH_2C(O)-$ or $-CH_2NHCH(R^{14})C(O)-$ wherein R^{14} is lower alkyl or (lower cycloalkyl)methyl; R^3 is absent or is $-N(R^{15})CH(R^{16})C(O)-$ wherein R^{15} is

hydrogen or methyl and R^{16} is hydrogen, lower alkyl, cyclopropylmethyl, cyclohexylmethyl, benzyl, $-\text{CH}(\text{OH})\text{CH}_3$, or $-(\text{CH}_2)_n\text{C}(\text{O})\text{OR}^{1*}$ or $-(\text{CH}_2)_n\text{C}(\text{O})\text{NR}^{11}\text{R}^{12}$ wherein n is the integer one, two or three, R^{1*} is hydrogen or a straight or branched chain alkyl containing one to ten carbon atoms, and R^{11} and R^{12} each independently is hydrogen, methyl or ethyl; and R^4 either is $-\text{NH}(\text{R}^{17})\text{CH}(\text{R}^{18})\text{C}(\text{O})-\text{Z}$ wherein R^{17} is hydrogen or methyl, R^{18} is lower alkyl, lower alkyl monosubstituted with a hydroxy or benzyloxy, cyclopropylmethyl, cyclohexylmethyl, (4-hydroxyphenyl)-methyl, $-\text{CH}(\text{OH})\text{C}_6\text{H}_5$ or benzyl, and Z is hydroxy or $\text{NR}^{19}\text{R}^{2*}$ wherein R^{19} and R^{2*} each independently is hydrogen, methyl, ethyl or 2-methylbutyl, or R^{19} and R^{2*} together with the nitrogen atom to which they are attached form a pyrrolidino or piperidino, or R^4 is $-\text{NR}^{17}\text{CR}^{18}-(\text{R}^{21})\text{CH}_2\text{OH}$ wherein R^{17} and R^{18} are as defined in the last instance; and R^{21} is hydrogen, lower alkyl or lower alkyl substituted with a hydroxy; or a therapeutically acceptable salt thereof.

Using the conventional three letter system for designating amino acid residues (see the second paragraph of the Details of the Invention, below), a more preferred group of the peptides for inhibiting HIV protease is represented by formula 1 wherein R^1 is the radical of formula 2 wherein R^5 is hydrogen, amino, nitro or $-\text{CH}_2\text{C}(\text{O})\text{OR}^7$ wherein R^7 is hydrogen, methyl, ethyl or benzyl, and L is hydrogen, R^2 is Val, Ala, Leu, Ile, Gly, Tbg, Cpa, Cha, Glu, Gln, Asn or Phe; Y is $-\text{NHCH}(\text{R}^{13})-\text{W}-$ wherein R^{13} is 1-methylethyl, 2-methylpropyl, cyclohexylmethyl, $-\text{CH}_2\text{CH}_2\text{SCH}_3$, benzyl or (4-methoxyphenyl)methyl, and W is $-\text{CH}(\text{OH})\text{CH}_2\text{C}(\text{O})-$ or $-\text{CH}_2\text{NHCH}(\text{R}^{14})\text{C}(\text{O})-$ wherein R^{14} is 2-methylpropyl or cyclohexylmethyl; R^3 is absent or is Leu, Nle, Ile, Val, Ala, Gly, Cha, Phe, Thr, Glu, Gln, Asp or Asn; R^4 is Leu- NH_2 , Leu-OH, Ile- NH_2 , Ile-OH, Val- NH_2 , Val-OH, Ala- NH_2 , Ala-OH, Thr(OBzl)- NH_2 , Cpa- NH_2 , Cpa-OH, Cha- NH_2 , Cha-OH, Tyr- NH_2 , Tyr-OH, Phe- NH_2 , Phe- $\text{NH}[\text{CH}_2\text{CH}(\text{CH}_3)\text{C}_2\text{H}_5]$, Phe- $\text{N}(\text{C}_2\text{H}_5)_2$, or Phe-OH, or R^4 is $-\text{NHCR}^{18}(\text{R}^{21})-\text{CH}_2\text{OH}$ wherein R^{18} is lower alkyl, $-\text{CH}_2\text{OH}$, $-\text{CH}(\text{OH})\text{CH}_3$, $\text{CH}_2\text{CH}_2\text{OH}$ or $-\text{CH}(\text{OH})\text{C}_6\text{H}_5$ and R^{21} is hydrogen, methyl or $-\text{CH}_2\text{OH}$; or a therapeutically acceptable salt thereof.

A most preferred group of peptides for inhibiting HIV protease is represented by compounds of formula 1 in which R^1 is 3-(1,2,3,4-tetrahydroisoquinolyl)carbonyl, R^2 and Y are as defined in the last instance; R^3 is Leu, Nle, Ile, Val, Ala, Cha, Glu or Gln, R^4 either is Ile- NH_2 , Ala- NH_2 , Thr(OBzl)- NH_2 , Cha- NH_2 or Phe- NH_2 or is $-\text{NHCR}^{18}(\text{R}^{21})\text{CH}_2\text{OH}$ wherein R^{18} is 1-methylethyl 1-methylpropyl or 2-methylpropyl, $-\text{CH}_2\text{OH}$ or $-\text{CH}(\text{OH})\text{C}_6\text{H}_5$ and R^{21} is hydrogen or methyl; or a therapeutically acceptable salt thereof.

Included within the scope of this invention is a pharmaceutical composition for treating HIV infections in a human, or for treating renin-dependent hypertension, comprising a compound of formula 1, or a therapeutically acceptable salt thereof, and a pharmaceutically acceptable carrier.

The scope of the invention includes as well a method for treating HIV infections in a human comprising administering thereto an effective amount of the compound of formula 1, or a therapeutically acceptable salt thereof.

Also included within the scope is a method for protecting human cells against HIV pathogenesis comprising treating said cells with an anti-HIV effective amount of a compound of formula 1, or a therapeutically acceptable salt thereof.

Still also included is a method of treating renin-associated hypertension or congestive heart failure in a mammal comprising administering thereto a renin-lowering effective amount of a compound of formula 1, or a therapeutically acceptable salt thereof.

Processes for preparing the compounds of formula 1 are described hereinafter.

Details of the Invention

GENERAL

The term "residue" with reference to an amino acid means a radical derived from the corresponding amino acid by eliminating the hydroxyl of the carboxy group and one hydrogen of the α -amino group.

In general, the abbreviations used herein for designating the amino acids and the protective groups are based on recommendations of the IUPAC-IUB Commission of Biochemical Nomenclature, see European Journal of Biochemistry, 138,9(1984). For instance, Val, Glu, Gln, Ala, Ile, Asp, Phe, Leu, Asn and Gly represent the residues of L-valine, L-glutamic acid, L-glutamine, L-alanine, L-isoleucine, L-aspartic acid, L-phenylalanine, L-leucine, L-asparagine and glycine, respectively. The symbols "Cpa" and "Cha" represent the residues of 2(S)-amino-3-cyclopropylpropionic acid (L-cyclopropylalanine) and 2(S)-amino-3-cyclohexylpropionic acid (L-cyclohexylalanine), respectively. The symbols "Nle" and "Tbg" represent the residues of 2(S)-aminohexanoic acid (1-norleucine) and 2(S)-amino-3,3-dimethylbutyric acid, respectively.

The term "lower alkyl" as used herein, either alone or in combination with a radical, means straight

chain alkyl radicals containing one to four carbon atoms and branched chain alkyl radicals containing three to four carbon atoms and includes methyl, ethyl, propyl, butyl, 1-methylethyl, 1-methylpropyl, 2-methylpropyl and 1,1-dimethylethyl.

The term "lower cycloalkyl" as used herein, either alone or in combination with a radical, means saturated cyclic hydrocarbon radicals containing from three to six carbon atoms and includes cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl.

The term "lower alkoxy" as used herein means straight chain alkoxy radicals containing one to four carbon atoms and branched chain alkoxy radicals containing three to four carbon atoms and includes methoxy, ethoxy, propoxy, 1-methylethoxy, butoxy and 1,1-dimethylethoxy. The latter radical is known commonly as tertiarybutyloxy.

The symbol "Boc" represents 1,1-dimethylethoxycarbonyl, known commonly as tertiary-butyloxycarbonyl. The symbol "C₆H₅" represents a phenyl radical.

The term "halo" as used herein means a halo radical selected from bromo, chloro, fluoro or iodo.

With reference to Y of general formula 1, the radical "-NHCH(R¹³)-W-" wherein R¹³ is as defined hereinabove and W is -CH(OH)CH₂CO- represents the radical derived from the amino acid known as statine (i.e. 4(S)-amino-3(S)-hydroxy-6-methylheptanoic acid) and its close analogs. The radical is derived by eliminating the hydroxyl of the carboxy group and one hydrogen of the amino group of the corresponding γ -amino acid. Each such radical has two chiral centers and thus can exist in various optically active or optically inactive forms. All forms are included for the peptides of formula 1 and for the appropriate intermediates therefore, the 4(S)-amino-3(S)-hydroxy enantiomers being preferred. The requisite 4-amino-3-hydroxy pentanoic acids for preparing the synthon to incorporate the radical into the peptide of formula 1 can be prepared by methods described by D.H. Rich and E.T.O. Sun, J. Med. Chem., 23, 27 (1980), and references therein.

The term "Sta" represents the radical -NHCH(2-methylpropyl)CH(OH)CH₂C(O)-, derived from statine. The term "ACHPA" represents the radical NHCH(cyclohexylmethyl)CH(OH)CH₂C(O)- derived from 4-amino-5-cyclohexyl-3-hydroxypentanoic acid, and the term "AHPPA" represents the radical -NHCH(benzyl)CH(OH)CH₂C(O)-, derived from 4-amino-5-phenyl-3-hydroxypentanoic acid. The 4(S)-amino-3(S)-hydroxy enantiomers of these last three embodiments are preferred. Unless designated otherwise by an antecedent such as (3R, 4R), the terms Sta, ACHPA and AHPPA represent their respective 4(S)-amino-3(S)-hydroxy enantiomers.

The asymmetric carbon atoms of the R¹, R², R³ and R⁴ units of the peptides of formula 1 have an S configuration, except those residing in the side chain of the amino acid or derived amino acid residues which may have the R configuration.

The term "amino" as used herein means an amino radical of formula -NH₂. The term "lower alkylamino" as used herein means alkylamino radicals containing one to six carbon atoms and includes methylamino, ethylamino, propylamino, 1-methyl-ethylamino and 2-methylbutylamino. The term "di(lower alkyl)amino" means an amino radical having two lower alkyl substituents each of which contains one to six carbon atoms and includes dimethylamino, diethylamino, ethylmethylamino and the like.

Additional abbreviations or symbols used hereafter for derived amino acid residues include THIQ for the R¹ radical 3-(1,2,3,4-tetrahydroisoquinolyl)carbonyl. Also note that when Y is the radical -NHCH(R¹³)-W- wherein R¹³ is as defined hereinabove and W is CH₂NHCH(R¹⁴)CO- wherein R¹⁴ is as defined hereinabove, the radical is equivalent to two adjoining, corresponding amino acid residues wherein the amide bond joining the two residues is reduced. According to convention, the latter radical can be expressed symbolically as two amino acid residues (in the three letter system) with the symbol " Ψ [CH₂NH]" inserted between the designation of the two adjoining amino acid residues. Accordingly, for example, the peptide of formula 1 wherein R¹ is 3-(1,2,3,4-tetrahydroisoquinolyl)carbonyl, R² is Val, Y is Sta, R³ is Leu, R⁴ is Phe-NH₂ is designated as THIQ-Val-Sta-Leu-Phe-NH₂; and the peptide of formula 1 wherein R¹ is 3-(1,2,3,4-tetrahydroisoquinolyl)carbonyl, R² is Ala, Y is -NHCH(benzyl)CH₂NHCH(2-methylpropyl) with two (S)-asymmetric centers, R³ is Ala, R⁴ is Phe-OH is designated as THIQ-Ala-Phe Ψ [CH₂NH]Leu-Ala-Phe-OH.

The term "coupling agent" as used herein means an agent capable of effecting the dehydrative coupling of an amino acid or peptide free carboxy group with a free amino group of another amino acid or peptide to form an amide bond between the reactants. The agents promote or facilitate the dehydrative coupling by activating the carboxy group. Descriptions of such coupling agents and activated groups are included in general textbooks of peptide chemistry; for instance, E. Schroder and K.L. Lubke, "The Peptides", Vol. 1, Academic Press, New York, N.Y., 1965, pp 2-128, and K.D. Kopple, "Peptides and Amino acids", W.A. Benjamin, Inc., New York, N.Y., 1966, pp 33-51. Examples of coupling agents are thionyl chloride, diphenylphosphoryl azide, dicyclohexylcarbodiimide, N-hydroxysuccinimide, or 1-hydroxybenzotriazole in the presence of dicyclohexylcarbodiimide. A very practical and useful coupling agent is

(benzotriazol-1-yloxy)tris(dimethylamino)phosphonium hexafluorophosphate, described by B. Castro et al., Tetrahedron Letters, 1219 (1975), see also D. Hudson, J. Org. Chem., 53, 617 (1988), either by itself or in the presence of 1-hydroxybenzotriazole.

The term "pharmaceutically acceptable carrier" as used herein means a non-toxic, generally inert vehicle for the active ingredient, which does not adversely affect the ingredient.

The term "effective amount" as used herein means a pre-determined amount of the peptide of this invention sufficient to be effective against HIV in vivo or renin-associated hypertension.

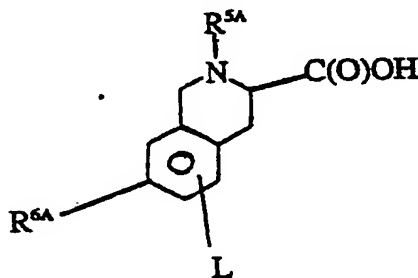
10 PROCESS

The peptides of formula 1 can be prepared by processes which incorporate therein methods commonly used in peptide synthesis such as classical solution coupling of amino acid residues and/or peptide fragments, and if desired solid phase techniques. Such methods are described, for example, by E. Schroder and K. Lubke, cited above, in the textbook series, "The Peptides: Analysis, Synthesis, Biology", E. Gross et al., Eds., Academic Press, New York, N.Y., 1979-1987, Volumes 1 to 8, and by J.M. Stewart and J.D. Young in "Solid Phase Peptide Synthesis", 2nd ed., Pierce Chem. Co., Rockford IL, USA, 1984.

A common feature of the aforementioned processes for the peptides is the protection of the labile side chain groups of the various amino acid residues or derived amino acid residues with suitable protective groups which will prevent a chemical reaction from occurring at that site until the protective group is ultimately removed. Usually also common is the protection of an α -amino group on an amino acid or a fragment while that entity reacts at the carboxy group, followed by the selective removal of the α -amino protective group to allow subsequent reaction to take place at that location. Usually another common feature is the initial protection of the C-terminal carboxyl of the amino acid residue or peptide fragment, if present, which is to become the C-terminal function of the peptide, with a suitable protective group which will prevent a chemical reaction from occurring at that site until the protective group is removed after the desired sequence of the peptide has been assembled.

Another feature of the preparation of the peptides of formula 1 is the incorporation into the peptide of the unit Y. As noted previously the unit Y can be either of two subunits, i.e. $-\text{NHCH}(\text{R}^{13})\text{CH}(\text{OH})\text{CH}_2\text{C}(\text{O})-$ wherein R^{13} is as defined herein or $-\text{NHCH}(\text{R}^{13})\text{CH}_2\text{NHCH}(\text{R}^{14})\text{C}(\text{O})-$ wherein R^{13} and R^{14} are as defined herein. The first-mentioned subunit can readily be incorporated into the peptide by coupling the N-protected derivative of the corresponding 4-amino-3-hydroxypentanoic acid (of formula $\text{NH}_2\text{CH}(\text{R}^{13})\text{CH}(\text{OH})\text{CH}_2\text{C}(\text{O})\text{OH}$), noted above, at the appropriate point during the preparation of the peptide by the classical methods of coupling of amino acid residues or fragments. The second-mentioned subunit, $-\text{NHCH}(\text{R}^{13})\text{CH}_2\text{NHCH}(\text{R}^{14})\text{C}(\text{O})-$, can be incorporated by forming the linear peptidyl framework of the peptide of formula 1, or a fragment thereof, by a reductive alkylation between two sub-fragments, each sub-fragment containing a precursor portion of the Y unit and at least one of the sub-fragments containing one or more of the amino acid units, whereby the CH_2NH bond of the Y unit is formed; for example, the reductive N-alkylation of the dipeptide of formula $\text{NH}_2\text{CH}(\text{R}^{14})\text{C}(\text{O})\text{R}^3\text{R}^4$ with $\text{Boc-NHCH}(\text{R}^{13})\text{CHO}$ in the presence of sodium cyanoborohydride to give the fragment $\text{Boc-NHCH}(\text{R}^{13})\text{CH}_2\text{NHCH}(\text{R}^{14})\text{C}(\text{O})\text{R}^3\text{R}^4$.

With reference to the process products, the peptides of formula 1 can be prepared by: coupling the derived amino acid of formula 3:

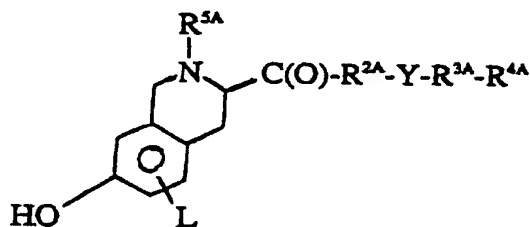


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wherein R^{5A} is lower alkyl, or $\text{QOC}(\text{O})-$ or $\text{QC}(\text{O})-$ wherein Q is as defined hereinabove; R^{6A} is hydrogen, hydroxy, lower alkyl, lower alkoxy, amino, nitro or $-\text{OCH}_2\text{C}(\text{O})\text{OR}^{7A}$ wherein R^{7A} is lower alkyl, lower cycloalkyl, (lower cycloalkyl)methyl, phenyl, phenyl monosubstituted with lower alkyl, lower alkoxy or halo,

benzyl or benzyl monosubstituted with lower alkyl, lower alkoxy or halo, and L is as defined hereinabove, with a fragment of formula $H-R^{2A}-Y-R^{3A}-R^{4A}$ wherein R^{2A} , R^{3A} and R^{4A} have the same meaning as defined herein for R^2 , R^3 and R^4 respectively, except that side chain amino and carboxy groups, and a C-terminal carboxy group, if present, are replaced with corresponding protected groups, and Y is as defined hereinabove; followed, if required, by amino deprotection and/or carboxy deprotection, to give the corresponding peptide of formula 1. Note that benzyl or substituted benzyl of R^{7A} can serve a dual role, i.e. serve as the progenitor for the corresponding radical in the ultimate product of the synthesis or serve as a carboxy protecting group. Also note that the radicals $QOC(O)-$ and $QC(O)-$ can serve a dual role as a progenitor for the corresponding radical in the final product or serve as an amino protecting group. When required, such carboxy and amino-protecting groups can be removed by known methods.

Alternatively, the peptides of formula 1 in which R^1 is the derived amino acid radical of formula 2 in which R^6 is $-OCH_2C(O)OR^7$ wherein R^7 is as defined herein and L, R^2 , Y, R^3 and R^4 are as defined herein can be prepared by subjecting the intermediate of formula 4



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wherein R^{2A} , Y, R^{3A} , R^{4A} and R^{5A} are as described herein to above, to O-alkylation with an alkylating agent of formula $XCH_2C(O)OR^{7A}$ wherein X is bromo, chloro or iodo and R^{7A} is as defined hereinabove, in the presence of a suitable strong base; followed, if required, by deprotection to obtain the desired corresponding peptide of formula 1. Suitable strong bases for the above-noted alkylation include alkali metal carbonates, preferably potassium carbonate; alkali metal hydroxides, preferably sodium hydroxide or potassium hydroxide; or alkali metal hydrides, preferably sodium hydride.

The derived amino acid of formula 3 can be prepared by known methods for preparing derivatives of 7-substituted-1,2,3,4-tetrahydro-3-isoquinolinecarboxylic acids; see G. Jones in "Comprehensive Chemistry", Vol 2, 1st ed., A.R. Katritzky and C.W. Rees, Eds, Pergamon Press, Oxford, UK, 1984, p 395. In turn, the derived amino acid of formula 3 in which R^{6A} is hydroxy is used to prepare the intermediate of formula 4 by conventional stepwise coupling procedures.

The peptide of formula 1 of this invention can be obtained in the form of a therapeutically acceptable salt.

In the instance where a particular peptide has a residue which functions as a base, examples of such salts are those with organic acids, e.g. acetic, lactic, succinic, benzoic, salicylic, methanesulfonic or p-toluenesulfonic acid, as well as polymeric acids such as tannic acid or carboxymethyl cellulose, and also salts with inorganic acids such as hydrohalic acids, e.g. hydrochloric acid, or sulfuric acid, or phosphoric acid if desired, a particular acid addition salt is converted into another acid addition salt, such as a non-toxic, pharmaceutically acceptable salt, by treatment with the appropriate ion exchange resin in the manner described by R.A. Boissonnas et al., Helv. Chim. Acta, 43, 1849 (1960).

In the instance where a particular peptide has one or more free carboxy groups, examples of such salts are those with the sodium, potassium or calcium cations, or with strong organic bases, for example, triethylamine or N-methylmorpholine.

In general, the therapeutically acceptable salts of the peptides of formula 1 are biologically fully equivalent to the peptides themselves.

BIOLOGICAL ASPECTS

The HIV protease inhibiting properties and the cell protective effect against HIV pathogenesis of the peptides of formula 1, or a therapeutically acceptable salt thereof, can be demonstrated by biochemical, microbiological and biological procedures.

A particular useful procedure for demonstrating the HIV protease inhibiting properties of the peptides of formula 1 or their therapeutically acceptable salts is the "Recombinant HIV Protease HLPC Assay". The procedure is based on the capacity of the test compound to inhibit enzymatic cleavage by HIV protease of

a decapeptide (the substrate) having an amino acid sequence which includes a known HIV protease cleavage site of the HIV poly-protein; see H.G. Krausslich et al., Proc. Natl. Acad. Sci. USA, 86, 807 (1989). Details of this assay together with the results obtained for exemplified peptides of formula 1 are described in the examples hereinafter.

5 The cell protective effect of the peptides or their therapeutically acceptable salts can be demonstrated by microbiological procedures for evaluating the effect of test compounds in inhibiting the cytopathogenicity of HIV in human T4 cell lines; for example, see M. Baba et al., Biochem. Biophys. Res. Comm., 142, 128 (1987).

10 When a peptide of this invention, or a therapeutically acceptable salt thereof, is used to combat HIV infections in a human, the peptide can be administered orally, topically or parenterally, in a vehicle comprising one or more pharmaceutically acceptable carriers, the proportion of which is determined by the solubility and chemical nature of the peptide, chosen route of administration and standard biological practice. For oral administration, the peptide or a therapeutically acceptable salt thereof can be formulated in unit dosage forms such as capsules or tablets each containing a predetermined amount of the active
15 ingredient, ranging from about 25 to 500 mg, in a pharmaceutically acceptable carrier. For topical administration, the peptide can be formulated in a pharmaceutically acceptable vehicle containing 0.1 to 10 percent, preferably 0.5 to 5 percent, of the active agent. Such formulations can be in the form of a cream, lotion, sublingual tablet, or preferably a transdermal patch or buccal patch.

For parenteral administration, the peptide of formula 1 is administered by either intravenous, subcutaneous or intramuscular injection, in compositions with pharmaceutically acceptable vehicles or carriers. For administration by injection, it is preferred to use the peptide in solution in a sterile aqueous vehicle which may also contain other solutes such as buffers or preservatives as well as sufficient quantities of pharmaceutically acceptable salts or of glucose to make the solution isotonic.

Suitable vehicles or carriers for the above noted formulations can be found in standard pharmaceutical texts, e.g. in "Remington's Pharmaceutical Sciences", 16th ed, Mack Publishing Company, Easton, Penn.,
25 1980.

The dosage of the peptide will vary with the form of administration and the particular active agent chosen. Furthermore, it will vary with the particular host under treatment. Generally, treatment is initiated with small dosages substantially less than the optimum dose of the peptide. Thereafter, the dosage is
30 increased by small increments until the optimum effect under the circumstances is reached. In general, the peptide is most desirably administered at a concentration level that will generally afford antivirally effective results without causing any harmful or deleterious side effects.

For oral administration, the peptide or a therapeutically acceptable salt is administered in the range of 1.0 to 75 mg per kilogram of body weight per day, with a preferred range of 2.5 to 20 mg per kilogram.

35 With reference to systemic administration, the peptide of formula 1 is administered at a dosage of 10 mcg to 1000 mcg per kilogram of body weight per day, although the aforementioned variations will occur. However, a dosage level that is in the range of from about 50 mcg to 500 mcg per kilogram of body weight per day is most desirably employed in order to achieve effective results.

Although the formulations disclosed hereinabove are effective and relatively safe medications for
40 treating HIV infections, the possible concurrent administration of these formulations with other antiviral medications or agents to obtain beneficial results is not excluded. Such other antiviral medications or agents include soluble CD4, zidovudine, dideoxycytidine, phosphonoformate, ribavarin, antiviral interferons (e.g. α -interferon or interleukin-2) or aerosol pentamidine.

The peptides of formula 1 also possess the ability to inhibit renin activity. The renin inhibiting activity of
45 the compounds can be demonstrated in standard pharmacological tests such as those described by M.G. Bock et al., J. Med. Chem., 31, 1918 (1988). As such the peptides are indicated for the diagnosis, prophylaxis and treatment of renin-associated hypertension in mammals including humans. The peptides also can be used for treating congestive heart failure in mammals including humans. For the latter purposes or indications, the peptides can be formulated and administered in the same manner as described above,
50 but usually at higher dosages which can be determined conventionally by using well known pharmacological protocols.

A preferred group of peptides of formula 1 for inhibiting renin is represented by formula 1 wherein R¹ is the radical of formula 2 wherein R⁵ is hydrogen or QOC(O)- wherein Q is lower alkyl, R⁵ is hydrogen, lower alkoxy, amino or nitro and L is hydrogen; R² is -NHCH(R³)C(O)- wherein R³ is lower alkyl, lower alkyl
55 monosubstituted with hydroxy, benzyl, 4-imidazolylmethyl, 2-thienylmethyl or 2-thiazolylmethyl, Y is -NHCH(R¹³)-W- wherein R¹³ is as defined herein and W is CH(OH)CH₂C(O)-, R³ is absent or is -NHCH(R¹⁵)-C(O)- wherein R¹⁵ is lower alkyl, and R⁴ is -NHCHR¹⁸C(O)-Z wherein R¹⁸ is lower alkyl, lower alkoxy monosubstituted with hydroxy or benzyloxy, -CH(OH)C₆H₅ or benzyl and Z is hydroxy or amino, or R⁴ is

-NHCR¹⁸(R²¹)CH₂OH wherein R¹⁸ is lower alkyl, lower alkyl monosubstituted with hydroxy, -CH(OH)C₆H₅ or benzyl, and R²¹ is hydrogen or methyl, or a therapeutically acceptable salt thereof.

A more preferred group of peptides for inhibiting renin is represented by formula 1 wherein R¹ is 3-(1,2,3,4-tetrahydroisoquinolyl)carbonyl or N-(tertiarybutyloxycarbonyl)-3-(1,2,3,4-tetrahydroisoquinolyl)-carbonyl, R² is Val, Ala or Phe, Y is Sta, ACHPA or AHPPA, R³ is absent or is Ala, Ile or Leu and R⁴ is Ala-NH₂, Phe-NH₂, Thr(OBzl)-NH₂ or -NHCR¹⁸(R¹⁹)CH₂OH wherein R¹⁸ is CH₂OH, -CH(OH)CH₃ or -CH(OH)-C₆H₅ and R¹⁹ is hydrogen or methyl, or a therapeutically acceptable salt thereof.

The following examples illustrate further this invention. Solution percentages or ratios express volume to volume relationship, unless stated otherwise. Abbreviations used in the examples include Boc: t-butyloxycarbonyl; BOP: (benzotriazol-1-yloxy)tris-(dimethylamino)-phosphonium hexafluorophosphate; Bzl: benzyl; DMF: dimethyl formamide; Et₂O diethyl ether; Fm: 9-fluorenylmethyl; HPLC: high performance liquid chromatography; MeOH: methanol; TFA: trifluoroacetic acid; THF: tetrahydrofuran.

Example 1

(Example of a procedure involving the stepwise coupling of units in sequence.)

Preparation of N-[N-[N-[3-(1,2,3,4-Tetrahydroisoquinolyl)carbonyl]-L-alanyl]-4(S)-amino-3(S)-hydroxy-5-cyclohexyl-pentanoyl]-L-leucyl]-L-phenylalaninamide Hydrochloride (THIQ-AlaACHPA-Leu-Phe-NH₂.HCl).

a) Boc-Phe-NH₂: Isobutyl chloroformate (2.86 mL, 22 mmol) was added dropwise at 0 °C to a stirred solution of N-methylmorpholine (2.42 mL, 22 mmol) and Boc-Phe-OH (5.31 g, 20 mmol) in dry THF. The resulting solution was stirred for an additional 30 min at 0 °C. Thereafter, a 28% aqueous solution of ammonia (5 mL) was added dropwise over 5 min. The solvent was evaporated and the residue was dissolved in ethyl acetate. The solution was washed successively with a 5% aqueous solution of citric acid (three times), saturated aqueous NaHCO₃ (three times) and saturated aqueous NaCl. The organic solution was dried over Na₂SO₄ and evaporated to afford Boc-Phe-NH₂ (5.0 g, 94%) as a white solid.

b) Boc-Leu-Phe-NH₂: A solution of Boc-Phe-NH₂ (6.46 g, 24.4 mmol) in 6N HCl/dioxane (84 mL) was stirred at room temperature (20-22 °C) under a nitrogen atmosphere for 30 min. The solvent was evaporated and the residue was dried under high vacuum. The solid residue was suspended in dry CH₃CN (300 mL) and the resulting mixture was cooled to 0-5 °C and stirred under a nitrogen atmosphere. Dry Et₃N (3.7 mL, 26 mmol) was added, followed by Boc-Leu-OH(monohydrate) (5.4 g, 22 mmol), BOP (10.69 g, 24.2 mmol) and more Et₃N (7.4 mL, 53 mmol). After 1.5 h, more BOP (4.28 g, 9.7 mmol) and Et₃N (1.48 mL, 10.6 mmol) were added. The reaction was stirred for an additional 15 min, then the CH₃CN was evaporated under reduced pressure and the residue was partitioned between a saturated aqueous solution of NaCl (200 mL) and ethyl acetate (3 times 200 mL). The combined organic solutions were washed successively with 100 mL each of a 10% aqueous solution of citric acid, water, a 5% aqueous solution of NaHCO₃ (three times) and water. The organic solution was dried over MgSO₄ and concentrated under reduced pressure. Chromatography of the residue over silica gel (eluent = ethyl acetate) gave a white gum which was triturated with Et₂O/hexane. The resulting solid was collected to afford Boc-Leu-Phe-NH₂ (7.7 g, 93%) as a white solid. Mass spectrum: 378 (M + H)⁺. Amino acid analysis: Leu, 1.00; Phe, 1.00.

c) Boc-ACHPA-Leu-Phe-NH₂: A solution of Boc-Leu-Phe-NH₂ (100 mg, 0.26 mmol) in 6N HCl/dioxane (1 mL) was stirred at room temperature under a nitrogen atmosphere for 15 min. The solvent was evaporated and the residue was dried under high vacuum for 1 h. The solid was suspended in dry CH₃CN (2 mL) and stirred under a nitrogen atmosphere. The solution was adjusted to pH 8 (wet pH paper) by the addition of N-methylmorpholine, then BocACHPA-OH (83 mg, 0.26 mmol) was added. The solution was stirred at room temperature for 1h (during which time pH 8 was maintained by the occasional addition of N-methylmorpholine). The mixture was poured into a saturated aqueous solution of NaCl. The aqueous solution was extracted twice with ethyl acetate. The combined organic extracts were washed successively with ice-cold 0.5 N aqueous HCl, 10% aqueous Na₂CO₃ (twice) and saturated aqueous NaCl (three times). The organic solution was dried over Na₂SO₄ and the solvent was evaporated. Chromatography of the residue over silica gel, eluting with 5% methanol in chloroform, afforded Boc-ACHPA-Leu-Phe-NH₂ (138 mg, 92%) as a white solid.

d) Boc-Ala-ACHPA-Leu-Phe-NH₂: A solution of Boc-ACHPA-Leu-Phe-NH₂ (200 mg, 0.35 mmol) in 6N

HCl/dioxane (2 mL) was stirred at room temperature under a nitrogen atmosphere for 20 min. The solvent was evaporated and the residue was dried under high vacuum for 1 h. The solid was suspended in dry CH₃CN (3 mL) and stirred under a nitrogen atmosphere. The solution was adjusted to pH 8 (wet pH paper) by the addition of N-methylmorpholine, then Boc-Ala-OH (74 mg, 0.39 mmol) was added, the pH was again adjusted to pH 8 (as before) and BOP (172 mg, 0.39 mmol) was added. The solution was stirred at room temperature for 1 h (during which time pH 8 was maintained by the occasional addition of N-methylmorpholine). The mixture was poured into a saturated aqueous solution of NaCl. The aqueous solution was extracted twice with ethyl acetate. The combined organic extracts were washed successively with ice-cold 0.5 N aqueous HCl, 10% aqueous Na₂CO₃ (twice) and saturated aqueous NaCl (three times). The organic solution was dried over Na₂SO₄ and the solvent was evaporated. Chromatography of the residue over silica gel, eluting with 5% methanol in chloroform, afforded Boc-Ala-ACHPA-Leu-Phe-NH₂ (178 mg, 79%) as a white solid.

e) Boc-THIQ-Ala-ACHPA-Leu-Phe-NH₂: A solution of Boc-Ala-ACHPA-Leu-Phe-NH₂ (110 mg, 0.17 mmol) in 6N HCl/dioxane (1 mL) was stirred at room temperature under a nitrogen atmosphere for 20 min. The solvent was evaporated and the solid residue was dried under high vacuum for 1 h. The solid was suspended in dry CH₃CN (2 mL). The mixture was stirred under a nitrogen atmosphere. The solution was adjusted to pH 8 (wet pH paper) by the addition of N-methylmorpholine, then Boc-THIQ-OH (53 mg, 0.19 mmol) was added, the pH was again adjusted to pH 8 (as before) and BOP (84 mg, 0.19 mmol) was added. The solution was stirred at room temperature for 1 h (during which time pH 8 was maintained by the occasional addition of N-methylmorpholine). The mixture was poured into a saturated aqueous solution of NaCl. The aqueous solution was extracted twice with ethyl acetate. The combined organic solutions were washed successively with ice-cold 0.5 N aqueous HCl, 10% aqueous Na₂CO₃ (twice) and saturated aqueous NaCl (three times). The organic solution was dried over Na₂SO₄ and the solvent was evaporated. Chromatography of the residue over silica gel, eluting with 5% methanol in chloroform, afforded Boc-THIQ-Ala-ACHPA-Leu-Phe-NH₂ (106 mg, 77%) as a white solid. Mass spectrum: 805 (M+H)⁺. Amino acid analysis: Ala, 1.00; Leu, 1.01; Phe, 0.98; ACHPA, 1.29. [Boc-THIQ-OH was prepared from 1,2,3,4-tetrahydro-3-isoquinoline-3-carboxylic acid described by S. Archer, J. Org. Chem. 16, 430(1951).]

f) Title Compound of this Example: A solution of Boc-THIQ-Ala-ACHPA-Leu-Phe-NH₂ (55 mg, 0.068 mmol) in 6N HCl/dioxane (1.5 mL) was stirred at room temperature under a nitrogen atmosphere for 20 min. The solvent was evaporated and the residue was dried under high vacuum for 1 h. The solid was triturated with Et₂O and the resulting suspension was filtered. The collected precipitate was dried under high vacuum for 17 h at room temperature to afford the title compound (47 mg, 93%) as a white solid. Mass spectrum: 705 (M-Cl)⁺. Amino acid analysis: Ala, 1.00; Leu, 1.01; Phe, 0.99; ACHPA, 1.16.

Example 2

THIQ-Leu-ACHPA-Leu-Phe-NH₂.HCl

The title compound was prepared by following the procedure of example 1 but substituting Boc-Leu-OH for Boc-Ala-OH. Mass spectrum: 747 (M-Cl)⁺. Amino acid analysis: Leu, 2.00; Phe, 0.98; ACHPA, 1.13.

Example 3

THIQ-Phe-ACHPA-Leu-Phe-NH₂.HCl

The title compound was prepared by following the procedure of example 1 but substituting Boc-Phe-OH for Boc-Ala-OH. Mass spectrum: 781 (M-Cl)⁺. Amino acid analysis: Leu, 0.97; Phe, 2.03, ACHPA, 1.12.

Example 4

THIQ-Val-ACHPA-Leu-Phe-NH₂.HCl

The title compound was prepared by following the procedure of example 1 but substituting Boc-Val-OH for Boc-Ala-OH. Mass spectrum: 733 (M-Cl)⁺. Amino acid analysis: Leu, 1.01; Phe, 1.00; Val, 0.99; ACHPA, 0.96.

Example 5THIQ-Cpa-ACHPA-Leu-Phe-NH₂.HCl

The title compound was prepared by following the procedure for example 1 but substituting Boc-Cpa-OH for Boc-Ala-OH. Mass spectrum 745 (M-Cl)⁺. Amino acid analysis: Leu, 1.01; Phe, 0.99; ACHPA, 0.96; Cpa, 0.98.

Example 6THIQ-Ala-ACHPA-Ala-Phe-NH₂.HCl

The title compound was prepared by following the procedure for example 1 but substituting Boc-Ala-OH for Boc-Leu-OH. Mass spectra: 663 (M-Cl)⁺. Amino acid analysis: Ala, 2.02; Phe, 0.98; ACHPA, 1.08.

Example 7

(Example of a procedure involving the coupling of fragments)

Preparation of THIQ-Val-ACHPA-Ala-Phe-NH₂.HCl

a) Boc-ACHPA-OMe: A solution of diazomethane in Et₂O was added portionwise to a vigorously stirred solution of Boc-ACHPA-OH (2.00g, 6.34 mmol) in Et₂O (20 mL) until a persistent yellow colour was obtained. Excess diazomethane was removed by passing a stream of dry nitrogen through the solution. Evaporation of the solvent gave a clear oil which crystallized on standing to afford Boc-ACHPA-OMe (2.16 g, 100%) as a white solid. NMR (200 MHz, CDCl₃): δ 0.70-1.90 (m, 22H including at δ 1.45 (s, 9H)), 2.55 (m, 2H), 3.30 (broad s, 1H), 3.65 (m, 1H), 3.71 (s, 3H), 4.20 (m, 1H), 4.70 (d, 1H).

b) Boc-Val-ACHPA-OMe: A solution of Boc-ACHPA-OMe (2.16 g, 6.56 mmol) in 6N HCl/dioxane (15 mL) was stirred at room temperature under a nitrogen atmosphere for 20 min. The solvent was evaporated and the residue was dried under high vacuum for 1 h. The solid was suspended in dry CH₃CN (20 mL) and the suspension was stirred under a nitrogen atmosphere. The resulting solution was adjusted to pH 8 (wet pH paper) by the addition of N-methylmorpholine. Boc-Val-OH (1.42 g, 6.56 mmol) was added to the solution. The pH was adjusted to pH 8 (as before) and BOP (2.90 g, 6.56 mmol) was added. The solution was stirred at room temperature for 1 h (during which time pH 8 was maintained by the occasional addition of N-methylmorpholine). The mixture was poured into saturated aqueous NaCl (50 mL). The aqueous solution was extracted twice with ethyl acetate. The combined organic solutions were washed successively with ice-cold 0.5 N aqueous HCl, 10% aqueous NaHCO₃ (twice) and saturated aqueous NaCl (three times). The organic solution was dried over Na₂SO₄ and the solvent was evaporated. Chromatography of the residue over silica gel, eluting with mixtures of increasing concentration of ethyl acetate in hexane (30-50%), afforded Boc-Val-ACHPA-OMe (2.44 g, 87%) as a white solid. Mass spectrum: 429 (M+H)⁺. NMR (200 MHz, CDCl₃): δ 0.70-1.90 (m, 28H, including at δ 0.90-1.05 (dd, 6H) and 1.45 (s, 9H)), 2.21 (m, 1H), 2.45 (m, 2H), 3.46 (broad s, 1H), 3.71 (s, 3H), 3.90 (m, 1H), 3.95-4.12 (m, 2H), 4.95 (broad d, 1H), 6.25

(broad d, 1H).

c) Boc-THIQ-Val-ACHPA-OMe: A solution of Boc-Val-CHPA-OMe (1.25 g, 292 mmol) in 6N HCl/dioxane (10mL) was stirred at room temperature under a nitrogen atmosphere for 20 min. The solvent was evaporated and the residue was dried under high vacuum for 1 h. The solid was suspended in dry CH₃CN (15 mL) and stirred under a nitrogen atmosphere. The resulting solution was adjusted to pH 8 (wet pH paper) by the addition of N-methylmorpholine. Boc-THIQ-OH (836 mg, 3.00 mmol) was added to the solution. The pH of the solution again was adjusted to pH 8 (as before) and BOP (1.33 g, 3.00 mmol) was added. The solution was stirred at room temperature for 1 h (during which time pH 8 was maintained by the occasional addition of N-methylmorpholine). The mixture was poured into saturated aqueous NaCl (30 mL). The aqueous solution was extracted twice with ethyl acetate. The combined organic solutions were washed successively with ice-cold 0.5 N aqueous HCl, 10% aqueous NaHCO₃ (twice) and saturated aqueous NaCl (three times). The organic solution was dried over Na₂SO₄ and the solvent was evaporated. Chromatography of the residue over silica gel, eluting with mixtures of increasing concentration of ethyl acetate in hexane (40-50%), afforded Boc-THIQ-Val-ACHPA-OMe (1.48 g, 86%) as a white solid. Mass spectrum: 588 (M+H)⁺.

d) Boc-THIQ-Val-ACHPA-OH: A 1 N aqueous solution of NaOH (2.27 mL, 2.27 mmol) was added dropwise over 30 min to a stirred solution of Boc-THIQ-Val-ACHPA-OMe (1.30 g, 2.27 mmol) in a mixture of methanol (10 mL) and water (2 mL). The resulting solution was stirred for one hour at room temperature. The methanol was evaporated. The residual aqueous solution was extracted once with ethyl acetate and then acidified to pH 3 (pH meter), first by the addition of solid citric acid and then finally by the addition of a 1 M aqueous solution of the same. The resulting suspension was extracted with ethyl acetate (twice). The combined organic extracts were washed with saturated aqueous NaCl (twice) and dried over Na₂SO₄. Evaporation of the solvent gave Boc-THIQ-Val-ACHPA-OH (1.20 g, 92%) as a white solid. Mass spectrum: 574(M+H)⁺.

e) Boc-Ala-Phe-NH₂: A solution of Boc-Phe-NH₂ (265 mg, 1.00 mmol) in excess 6N HCl/dioxane was stirred at room temperature under a nitrogen atmosphere for 15 min. The solvent was evaporated and the residue was suspended in dry CH₃CN (2 mL). The suspension was stirred under a nitrogen atmosphere. The solution was adjusted to pH>8 (wet pH paper) by the addition of excess N-methylmorpholine. Boc-Ala-OH (189 mg, 1.00 mmol) and BOP (486 mg, 1.10 mmol) were added. The solution was stirred at room temperature for 2 h then poured into a mixture of saturated aqueous NaCl and ethyl acetate. The organic solution was separated then washed successively with saturated aqueous NaHCO₃ (three times), 0.5 N aqueous HCl (three times) and saturated aqueous NaCl. The organic solution was dried over Na₂SO₄ and the solvent was evaporated. Chromatography of the residue over silica gel, eluting with 10% methanol in chloroform, afforded Boc-Ala-Phe-NH₂ (173 mg, 51%) as a white solid.

f) Boc-THIQ-Val-ACHPA-Ala-Phe-NH₂: A solution of Boc-Ala-Phe-NH₂ (33.6 mg, 0.10 mmol) in excess 6N HCl/dioxane was stirred at room temperature under a nitrogen atmosphere for 15 min. The solvent was evaporated. The residue was suspended in dry CH₃CN and stirred under a nitrogen atmosphere. The solution was adjusted to pH>8 (wet pH paper) by the addition of excess N-methylmorpholine, then Boc-THIQ-Val-ACHPA-OH (55.7 mg, 0.10 mmol) and BOP (48.6 mg, 0.11 mmol) were added. The solution was stirred at room temperature for 2 h then poured into a mixture of saturated aqueous NaCl and ethyl acetate. The organic solution was washed successively with 0.5 N aqueous HCl (three times), saturated aqueous NaHCO₃ (three times) and saturated aqueous NaCl. The organic solution was dried over Na₂SO₄ and the solvent was evaporated. Chromatography of the residue over silica gel, eluting with mixtures of increasing concentration of methanol in chloroform, afforded Boc-THIQ-Val-ACHPA-Ala-Phe-NH₂ (30 mg, 38%) as a white solid.

g) Title Compound of this Example: A solution of Boc-THIQ-Val-ACHPA-Ala-Phe-NH₂ (30 mg, 0.038 mmol) in excess 6 N HCl/dioxane was stirred at room temperature under a nitrogen atmosphere for 15 min. The solvent was evaporated and the residue was triturated with Et₂O. The resulting suspension was filtered. The collected precipitate was dried to afford the title compound (14 mg, 51%) as a white solid. Mass spectrum: 691 (M-Cl)⁺. Amino acid analysis: Ala, 1.01; Phe, 0.95; Val, 1.03; ACHPA, 1.00.

By following the procedure for example 7 but substituting the appropriate Boc amino acid for Boc-Ala-OH, the following peptides of formula 1 were obtained:

Example 8: THIQ-Val-ACHPA-Phe-Phe-NH₂.HCl, MS: 767 (M-Cl)⁺, amino acid analysis: Phe, 1.95; Val, 1.05; ACHPA, 1.14.

Example 9: THIQ-Val-ACHPA-Gly-Phe-NH₂.HCl, MS: 677 (M-Cl)⁺, amino acid analysis: Gly, 1.02; Phe, 0.96; Val, 1.02; ACHPA, 1.08.

Example 10: THIQ-Val-ACHPA-Asn-Phe-NH₂.HCl, MS: 734 (M-Cl)⁺, amino acid analysis: Asx, 1.03; Phe, 0.97; Val, 1.00; ACHPA, 1.02.

Example 11: THIQ-Val-ACHPA-Gln-Phe-NH₂.HCl, MS: 748 (M-Cl)⁺, amino acid analysis: Glx, 1.02; Phe, 0.96; Val, 1.02; ACHPA, 1.06.

Example 12

THIQ-Val-ACHPA-Asp-Phe-NH₂.HCl

By following the procedure for example 7 but substituting Boc-Asp(O-Bzl)-OH for Boc-Ala-OH, Boc-THIQ-Val-ACHPA-Asp(O-Bzl)-Phe-NH₂ was obtained. The latter compound (77.1 mg, 0.10 mmol) was dissolved in MeOH (10 ml). Under an atmosphere of nitrogen, 10% palladium on carbon (7.7 mg) was added to the solution. The mixture was shaken on a Parr apparatus under an atmosphere of H₂ (50 psi) for 30 min. The solution was filtered through a 45 μm membrane and the filtrate was concentrated. Chromatography of the residue over silica gel, eluting with mixtures of increasing concentration of methanol in chloroform, afforded a homogeneous oil. The oil was dissolved in excess 6 N HCl/dioxane. The solution was stirred at room temperature under a nitrogen atmosphere for 15 min. The solvent was evaporated and the residue was triturated with Et₂O. The resulting suspension was filtered. The collected material was dried to afford the title compound (37.7 mg, 49%) as a white solid. Mass spectrum: 735 (M-Cl)⁺. Amino acid analysis: Asx, 1.00; Phe, 1.00; Val, 1.00.

Example 13

THIQ-Val-ACHPA-Glu-Phe-NH₂.HCl

The title compound was prepared by following the procedure of example 12 but substituting Boc-Glu(O-Bzl)-OH for Boc-Asp(O-Bzl)-OH. Mass spectrum: 749 (M-Cl)⁺. Amino acid analysis: Glx, 1.04; Phe, 0.98; Val, 0.98.

Example 14

THIQ-Ala-ACHPA-Leu-Phe-OH.HCl

By following the procedure for example 1 but substituting Boc-Phe-OFm for Boc-Phe-NH₂, Boc-THIQ-Ala-ACHPA-Leu-Phe-OFm was obtained. The latter compound (249 mg, 0.3 mmol) was dissolved in DMF (5 mL). Under an atmosphere of nitrogen, piperidine (296 μL, 3 mmol) was added at 0 °C to the stirred solution. The reaction mixture was stirred for 2 h at room temperature then diluted with ethyl acetate (20 mL) and washed successively with a 10% aqueous solution of citric acid, water and a saturated aqueous solution of NaCl. The organic solution was dried over MgSO₄ and concentrated under reduced pressure. Chromatography of the residue over silica gel, eluting with a mixture of CHCl₃/methanol/water (90:7:3), gave Boc-THIQ-Ala-ACHPA-Leu-Phe-OH (161 mg, 66%). Deprotection of the latter compound according to the general deprotection procedure of example 1(f) gave the title compound. Mass spectrum: 706 (M-Cl)⁺. Amino acid analysis: Ala, 0.98; Leu, 1.00; Phe, 1.03; ACHPA, 1.04.

Example 15

THIQ-Ala-AHPPA-Leu-Phe-NH₂.HCl

The title compound was prepared following the procedure for example 1 but substituting Boc-AHPPA-OH for Boc-ACHPA-OH. Mass spectrum: 699 (MCl)⁺.

Example 16

THIQ-Val-AHPPA-Leu-Phe-NH₂.HCl

The title compound was prepared following the procedure for example 1 but substituting Boc-AHPPA-OH for Boc-ACHPA-OH and Boc-Val-OH for Boc-Ala-OH. Mass spectrum: 727 (M-Cl)⁺.

Example 17

THIQ-Ala-Sta-Leu-Phe-NH₂.HCl

The title compound was prepared following the procedure for example 1 but substituting Boc-Sta-OH for Boc-ACHPA-OH. Mass spectrum: 655 (M-Cl)⁺. Amino acid analysis: Ala, 1.00; Leu, 1.00; Phe, 1.00; Sta, 1.04.

Example 18

THIQ-Val(3R,4S)AHPPA-Leu-Phe-NH₂.HCl

The title compound was prepared following the procedure for example 1 but substituting Boc-(3R,4S)-AHPPA-OH for Boc-ACHPA-OH and Boc-Val-OH for Boc-Ala-OH. Mass spectrum: 727 (M-Cl)⁺.

Example 19

THIQ-Val-Pheψ[CH₂NH]Leu-Leu-Phe-NH₂.HCl

a) H-Leu-Leu-Phe-NH₂.HCl

By following the coupling and deprotection procedures of example 1, H-Leu-Leu-Phe-NH₂.HCl was obtained.

b) Boc-Pheψ[CH₂NH]Leu-Leu-NH₂

Sodium cyanoborohydride (35 mg, 0.56 mmol) was added portionwise during 30 min to a stirred solution of Boc-phenylalaninal (0.70 mmol), described by D.H. Rich and E.T.O. Sun, supra, and H-Leu-Leu-Phe-NH₂.HCl (0.5 mmol) in MeOH (3 mL). The reaction mixture was stirred for 18 h at room temperature then cooled in ice. A saturated aqueous solution of NaHCO₃ (3 mL) was added, followed by ethyl acetate (10 mL). The organic phase was separated, washed successively with water and a saturated aqueous solution of NaCl, then dried over MgSO₄ and concentrated under reduced pressure. Chromatography of the residue over silica gel, eluting with a 3% solution of methanol in chloroform, gave Boc-Pheψ[CH₂NH]Leu-Leu-NH₂ (174 mg, 56%).

Title compound of this example.

The title compound was prepared from Boc-Pheψ[CH₂NH]Leu-Leu-Phe-NH₂ by using the general coupling and deprotection procedures described in example 1. Mass spectrum: 782 (M-2HCl)⁺.

Example 20THIQ-Val-Chaψ[CH₂NH]Leu-Leu-Phe-NH₂.HCl

The title compound was prepared by following the procedures described for example 19 but substituting Boc-cyclohexylalaninal for Boc-phenylalaninal. Mass spectrum: 788 (M-HCl₂)⁺. By following the procedures for example 19 but substituting the appropriate Boc amino acids, followed by subsequent purification by HPLC, the following compounds of formula 1 were obtained.

Example 21: THIQ-Val-Pheψ[CH₂NH]Leu-Gln-Ile-NH₂.2TFA. Mass spectrum: 763 (M-2TFA + H)⁺.

Example 22: THIQ-Asn-Pheψ[CH₂NH]Leu-Gln-Ile-NH₂.2TFA. Mass spectrum: 778 (M-2TFA + H)⁺.

Other examples of peptides of formula 1 include:

THIQ-Ile-ACHPA-Cha-Phe-NH₂

THIQ-Val-ACHPA-Cha-Phe-NH₂

THIQ-Ile-ACHPA-Leu-Phe-NH₂

THIQ-Ala-ACHPA-ASP(O(CH₂)₇CH₃)Phe-NH₂

3(7-hydroxy-1,2,3,4-tetrahydroisoquinoly)carbonyl-Cpa-AHPPA-Leu-Phe-OH

THIQ-Phe-Sta-Cha-Phe-OEt

THIQ-Val-ACHPA-Ile-NHCH[CH(OH)C₆H₅]CH₂OH

THIQ-Val-ACHPA-Cha-NHCH[CH(CH₃)C₂H₅]CH₂OH

THIQ-Ile-AHPPA-Phe-NH₂

Example 23Recombinant HIV Protease HPLC Assay:

Enzyme: HIV protease was expressed in E. coli and purified to ca. 50% purity according to the procedure described by H.-G. Krausslich et al., Proc. Natl. Acad. Sci. USA, 86, 807 (1989). The enzyme was stored as 10 μL aliquots at -70 °C. The aliquots were diluted to 1/10th of the original concentration with buffer prior to use (enzyme working solution).

Substrate: VSFNFPQITL-NH₂, MW 1164, see Krausslich et al., supra, was used as substrate. The substrate was made into 10 mM stock in DMSO and stored at -20 °C. Prior to use, the stock was diluted with buffer to give a 400 μM solution substrate working solution.

Buffer: 2-(4-Morpholino)ethanesulfonic acid (50mM), NaCl (25mM) and EDTA (5mM) was dissolved in distilled H₂O (90mL) and the solution was adjusted to pH6 with concentrated aqueous NaOH. The latter solution was diluted to 100 mL with H₂O to give the buffer.

Procedure: (1) The test compound was dissolved in DMSO to give a solution having 40X the final concentration of the test compound in the assay mixture (see step 2) so that the amount of DMSO in the assay mixture was 5% (v,v) or less. (2) The assay mixture was prepared by mixing 10 μL of the substrate working solution, 0.5 μL of the solution of the test compound in DMSO from step 1, and 10 μL of the enzyme working solution. (3) The assay mixture was incubated at 37 °C for 1h. (4) The reaction was quenched by adding 100 μL of 2% aqueous TFA. (5) The substrate and products (i.e. VSFNF and PQITL-NH₂) were separated by subjecting 100 μL of the quenched assay mixture to HPLC using Nucleosil ® C₁₈ column with a 26-min linear gradient at 1 mL/min from 11% to 70% acetonitrile in H₂O with 0.05% aqueous TFA. Elution was monitored at 210 nm. (6) A control which was the assay mixture without the test compound, was subjected simultaneously to steps 3 to 5.

Inhibition Studies: Cleavage products and remaining parent substrate were quantified by either peak height or by integration of the appropriate HPLC peaks. Substrate conversion was calculated using the

following relationship:

$$\% \text{ Conversion} = \frac{\text{Sum of peak height or peak area of products}}{\text{Sum of peak height or peak area of substrate and products}} \times 100$$

Enzyme inhibition of the test compound was calculated as follows:

$$\% \text{ Inhibition} = 100 - \frac{\% \text{ Conversion for assay mixture}}{\% \text{ Conversion of control}} \times 100$$

The concentration of the test compound which causes a 50% inhibition of the HIV-protease, i.e. the IC_{50} , was determined as follows: The percent inhibition of the enzyme was determined for a minimum of three different concentrations of the test compound. Thereafter, the IC_{50} was determined graphically by plotting the percent inhibition of the substrate against the concentration of the test compound.

The following table of exemplified peptides of formula 1 lists their IC_{50} as determined in the recombinant HIV protease HPLC assay.

Peptide	Example in which peptide is prepared	IC_{50} (nM)
THIQ-Ala-ACHPA-Leu-Phe-NH ₂ .HCl	1	90
THIQ-Val-ACHPA-Ala-Phe-NH ₂ .HCl	7	17
THIQ-Ala-ACHPA-Leu-Phe-OH.HCl	14	200
THIQ-Val-AHPPA-Leu-Phe-NH ₂ .HCl	16	25
THIQ-Val-Cha [CH ₂ NH]Leu-Leu-Phe-NH ₂ .2HCl	25	80

Example 24

Boc-1,2,3,4-Tetrahydro-7-nitro-3-isoquinolinecarboxylic Acid (3a: W = Boc and R^{7A} = NO₂)

(S)-1,2,3,4-Tetrahydro-7-nitroisoquinoline ethyl ester (50.0 mg, 0.2 mmol), described by Y. Oka and K. Nishikawa, European patent application 18,104 published October 29, 1980, was dissolved in dioxane-H₂O (1:1.2 mL). Diisopropylamine (53 μ L, 0.3 mmol) was added to the solution followed by the addition of a solution of ditertiary-butyl dicarbonate (52 mg, 0.24 mmol) in dioxane (0.5 mL). After 3h, the reaction mixture was diluted with ethyl acetate (15 mL). The organic solution was washed with cold 1N aqueous HCl, saturated aqueous NaHCO₃ (twice) and saturated aqueous NaCl. The organic layer was dried (Na₂SO₄) and concentrated. Chromatography of the residue over silica gel, eluting with 25% ethyl acetate in hexane, gave Boc(S)-1,2,3,4-tetrahydro-7-nitro-3-isoquinolinecarboxylic acid ethyl ester (85.6 mg) as a colorless oil.

A solution of later compound (13.6 mg, 0.053 mmol) in MeOH (0.5 mL) and H₂O (0.3 mL) was mixed with 1N aqueous NaOH (64 μ L, 0.064 mmol). After 3h, the solution was diluted with H₂O (10 mL) and washed with ethyl acetate (3 times 4 mL). The aqueous phase was mixed with fresh ethyl acetate and the mixture acidified to pH 4 with citric acid. The organic layer was separated and the aqueous phase extracted with ethyl acetate. The combined organic layer and extracts were washed with saturated aqueous NaCl, dried (Na₂SO₄) and concentrated to give the title compound (16.1 mg, 94%) as a colorless oil. Mass spectrum: 323 (M + H)⁺.

Example 25

3-(1,2,3,4-Tetrahydro-7-nitroisoquinolyl)carbonyl-Val-ACHPA-Leu-Phe-NH₂.HCl

The title compound was prepared by following the procedure of example 4 but substituting Boc-1,2,3,4-tetrahydro-7-nitro-3-isoquinolinecarboxylic acid for Boc-THIQ-OH. Mass spectrum: 778 (M + H)⁺.

Example 26

3-(7-Amino-1,2,3,4-tetrahydroisoquinolyl)carbonyl-Val-ACHPA-Leu-Phe-NH₂.2HCl

The title compound was prepared by following the procedure of example 25 but substituting the di-Boc derivative of 7-amino-1,2,3,4-tetrahydro-3-isoquinolinecarboxylic acid (prepared via H₂/Pt hydrogenation of Boc-1,2,3,4-tetrahydro-7-nitro-3-isoquinolinecarboxylic acid ethyl ester) for Boc-1,2,3,4-tetrahydro-7-nitro-3-isoquinolinecarboxylic acid. Mass spectrum: 748 (M + H)⁺.

Example 27

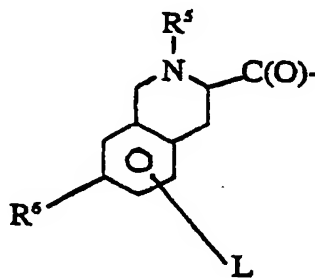
THIQ-Val-ACHPA-Cha-NH[CH(CH₃)C₂H₅]CH₂OH.HCl

The title compound is prepared by following the procedure of example 4 but substituting Boc-Cha-OH for Boc-Leu-OH and substituting Boc-isoleucinol for Boc-Phe-NH₂.

Claims

1. A peptide of formula 1

$R^1-R^2-Y-R^3-R^4$ ¹
wherein R¹ is a derived amino acid radical of formula 2



2

wherein R⁵ is hydrogen, lower alkyl, or QOC(O)- or QC(O)- wherein Q is lower alkyl, lower cycloalkyl, (lower cycloalkyl)methyl, phenyl, phenyl monosubstituted with lower alkyl, lower alkoxy or halo, benzyl or benzyl monosubstituted with lower alkyl, lower alkoxy or halo; R⁶ is hydrogen, hydroxy, lower alkyl, lower alkoxy, amino, nitro or -OCH₂H(O)OR⁷ wherein R⁷ is hydrogen, lower alkyl, lower cycloalkyl, (lower cycloalkyl)methyl, phenyl, phenyl monosubstituted with lower alkyl, lower alkoxy or halo, benzyl, or benzyl monosubstituted with lower alkyl, lower alkoxy or halo; and L on the aromatic ring of the radical of formula 2

represents hydrogen or a substituent on the aromatic ring; the substituent being selected from the group of lower alkyl, lower alkoxy and halo, or L represents the same or different of two substituents on the aromatic ring, the substituents being selected from the group of lower alkyl, lower alkoxy and halo, provided that any two substituents do not interfere with each others presence;

two substituents do not interfere with each others presence;
 5 R² is -N(R⁸)CH(R⁹)C(O)- wherein R⁸ is hydrogen or lower alkyl and R⁹ is hydrogen, lower alkyl, lower alkyl
 monosubstituted with hydroxy, methoxy, methylthio or benzyloxy, lower cycloalkyl, (lower cycloalkyl)methyl,
 benzyl, 4-imidazolylmethyl, 2-thiazolylmethyl, (4-hydroxyphenyl)methyl, [4-(lower alkoxy)phenyl]methyl, -
 CH(OH)C₆H₅, -(CH₂)₄NH₂, or -(CH₂)_nC(O)OR^{1*} or -(CH₂)_nC(O)NR¹¹R¹² wherein n is the integer one, two or
 10 three, R^{1*} is hydrogen, a straight or branched chain alkyl containing one to ten carbon atoms, or phenyl-
 (lower)alkyl and R¹¹ and R¹² each independently is hydrogen, lower alkyl, phenyl(lower)alkyl, or R¹¹ and R¹²
 together with the nitrogen to which they are joined form a pyrrolidino, piperidino, morpholino, piperazino or
 4-(lower alkyl)piperazino;

4-(lower alkyl)piperazino;
Y is a derived amino acid radical of the formula $-\text{NHCH}(\text{R}^{13})-\text{W}-$ wherein R^{13} is lower alkyl, (lower cycloalkyl)methyl, $-\text{CH}_2\text{CH}_2\text{SCH}_3$, benzyl or benzyl substituted on the aromatic portion thereof with hydroxy or lower alkoxy, and W is $-\text{CH}(\text{OH})\text{CH}_2\text{C}(\text{O})-$ or $-\text{CH}_2\text{NHCH}(\text{R}^{14})\text{C}(\text{O})-$ wherein R^{14} has the same meaning as defined for R^{13} ;

defined for R¹³;
R³ is absent or is -N(R¹⁵)CH(R¹⁶)C(O)- wherein R¹⁵ is hydrogen or lower alkyl and R¹⁶ has the same meaning as defined herein for R⁹; and
R⁴ is [-NR¹⁷CH(R¹⁸)C(O)]_p-Z wherein R¹⁷ is hydrogen or lower alkyl, R¹⁸ has the same meaning as defined herein for R⁹, p is the integer zero or one, and Z is hydroxy, lower alkoxy, benzyloxy or -NR¹⁹R^{2*} wherein R¹⁹ and R^{2*} each independently is hydrogen, lower alkyl or phenyl(lower)alkyl, or R¹⁹ and R^{2*} together with the nitrogen atom to which they are joined form a pyrrolidino, piperidino, morpholino, piperazino or 4-(lower alkyl)piperazino; or

25 (lower alkyl)piperazino; or
R⁴ is -NR¹⁷CR¹⁸(R²¹)CH₂OH wherein R¹⁷ and R¹⁸ are as defined in this claim and R²¹, is hydrogen, lower alkyl or lower alkyl monosubstituted with hydroxy; or a therapeutically acceptable salt thereof.

2. A peptide as recited in claim 1 wherein R¹ is the radical of formula 2 wherein R⁵ is hydrogen, R⁶ is hydrogen, hydroxy, methyl, methoxy, amino, nitro or -CH₂C(O)OR⁷ wherein R⁷ is hydrogen, lower alkyl or benzyl, and L is hydrogen or one or two halo substituents, R² is -N(R⁸)CH(R⁹)CO- wherein R⁸ is hydrogen or methyl and R⁹ is lower alkyl, lower cycloalkyl, cyclopropylmethyl, cyclohexylmethyl, -CH₂CH₂COOH, -CH₂CH₂CONH₂, -CH₂CONH₂ or benzyl; Y is -NHCH(R¹³)-W- wherein R¹³ is lower alkyl, (lower cycloalkyl)-methyl, -CH₂CH₂SCH₃, benzyl or (4-methoxyphenyl)methyl and W is -CH(OH)CH₂C(O)- or -CH₂NHCH(R¹⁴)-C(O)- wherein R¹⁴ is lower alkyl or (lower cycloalkyl)methyl; R³ is absent or is -N(R¹⁵)CH(R¹⁶)C(O)- wherein R¹⁵ is hydrogen or methyl and R¹⁶ is hydrogen, lower alkyl, cyclopropylmethyl, cyclohexylmethyl, benzyl, -CH(OH)CH₃, or (CH₂)_nC(O)OR¹⁰ or -(CH₂)_nC(O)NR¹¹R¹² wherein n is the integer one, two or three, R¹¹ is hydrogen or a straight or branched chain alkyl containing one to ten carbon atoms, and R¹² each independently is hydrogen, methyl or ethyl; and R⁴ either is -NH(R¹⁷)CH(R¹⁸)C(O)-Z wherein R¹⁷ is hydrogen or methyl, R¹⁸ is lower alkyl, lower alkyl monosubstituted with hydroxy or benzyloxy, cyclopropylmethyl, cyclohexylmethyl, (4-hydroxyphenyl)methyl, CH(OH)C₆H₅ or benzyl; and Z is hydroxy or NR¹⁹R²⁰ wherein R¹⁹ and R²⁰ each independently is hydrogen, methyl, ethyl or 2-methylbutyl or R¹⁹ and R²⁰ together with the nitrogen atom to which they are attached form a pyrrolidino or piperidino, or R⁴ is -NR¹⁷CR¹⁸(R²¹)CH₂OH wherein R¹⁷ and R¹⁸ are as defined in this claim and R²¹ is hydrogen, lower alkyl or lower alkyl monosubstituted with hydroxy; or a therapeutically acceptable salt thereof.

3. A peptide as recited in claim 2 wherein R¹ is the radical of formula 2 wherein R⁵ is hydrogen, R⁶ is hydrogen, amino, nitro or -CH₂C(O)OR⁷ wherein R⁷ is hydrogen, methyl, ethyl or benzyl, and L is hydrogen; R² is Val, Ala, Leu, Ile, Gly, Tbg, Cpa, Cha, Glu, Gln, Asn or Phe; Y is -NHCH(R¹³)-W- wherein R¹³ is 1-methylethyl, 2-methylpropyl, cyclohexylmethyl, -CH₂CH₂SCH₃, benzyl or (4-methoxyphenyl)methyl, and W is -CH(OH)CH₂C(O)- or CH₂NHCH(R¹⁴)C(O)- wherein R¹⁴ is 2-methylpropyl or cyclohexylmethyl; R³ is absent or is Leu, Nle, Ile, Val, Ala, Gly, Cha, Phe, Thr, Glu, Gln, Asp or Asn; and R⁴ is Leu-NH₂, Leu-OH, Ile-NH₂, Ile-OH, Val-NH₂, Val-OH, Ala-NH₂, Ala-OH, Thr(OBzl)-NH₂, Cpa-NH₂, Cpa-OH, Cha-NH₂, Cha-OH, Tyr-NH₂, Tyr-OH, Phe-NH₂, Phe-NH[CH₂CH(CH₃)C₂H₅], Phe-N(C₂H₅)₂ or Phe-OH, or R⁴ is -NHCR¹⁸(R²¹)-CH₂OH wherein R¹⁸ is lower alkyl, CH₂OH, -CH(OH)CH₃, CH₂CH₂OH or -CH(OH)C₂H₅ and R²¹ is hydrogen, methyl or CH₂OH; or a therapeutically acceptable salt thereof.

4. A peptide as recited in claim 3 wherein R¹ is 3-(1,2,3,4-tetrahydroisoquinolyl)carbonyl, R² and Y are as defined in claim 3, R³ is Leu, Ile, Val, Ala, Cha, Glu or Gln, R⁴ is either Ile-NH₂, Ala-NH₂, Thr(OBzl)-NH₂, Cha-NH₂ or Ph-NH₂, or is -NHCR¹⁸-(R²¹)CH₂OH wherein R¹⁸ is 1-methylethyl, 1-methylpropyl, 2-methylpropyl, -CH₂OH or -CH(OH)C₆H₅, and R²¹ is hydrogen or methyl; or a therapeutically acceptable salt thereof.

5. A peptide as recited in claim 1 wherein R¹ is the radical of formula 2 wherein R⁵ is hydrogen or

QOC(O)- wherein Q is lower alkyl, R⁶ is hydrogen, lower alkoxy, amino or nitro and L is hydrogen; R² is -NHCH(R³)C(O)- wherein R³ is lower alkyl, lower alkyl substituted with a hydroxy, benzyl, 4-imidazolylmethyl, 2-thienylmethyl or 2-thiazolylmethyl, Y is -NHCH(R¹³)-W- wherein R¹³ is as defined in claim 1 and W is -CH(OH)CH₂C(O)-, R³ is absent or is -NHCH(R¹⁶)C(O)- wherein R¹⁶ is lower alkyl, and R⁴ is NHCHR¹⁸C(O)-Z wherein R¹⁸ is lower alkyl, lower alkoxy monosubstituted with hydroxy or benzyloxy, -CH(OH)C₂H₅ or benzyl and Z is hydroxy or amino, or R⁴ is -NHCR¹⁸(R²¹)CH₂OH wherein R¹⁸ is lower alkyl, lower alkyl monosubstituted with hydroxy, CH(OH)C₆H₅ or benzyl, and R²¹ is hydrogen or methyl, or a therapeutically acceptable salt thereof.

6. A peptide as recited in claim 5 wherein R¹ is 3-(1,2,3,4-tetrahydroisoquinolyl)carbonyl or N-(tertiary-butyloxycarbonyl)-3-(1,2,3,4-tetrahydroisoquinolyl)carbonyl, R² is Val, Ala or Phe, Y is Sta, ACHPA or AHPPA, R³ is absent or is Ala, Ile or Leu and R⁴ is Ala-NH₂, Phe-NH₂, Thr(OBzl)-NH₂ or -NHCR¹⁸(R¹⁹)-CH₂OH wherein R¹⁸ is -CH₂OH, -CH(OH)CH₃ or -CH(OH)C₆H₅ and R¹⁹ is hydrogen or methyl, or a therapeutically acceptable salt thereof.

7. A peptide as recited in claim 1 selected from the group consisting of:

- 15 THIQ-Ala-ACHPA-Leu-Phe-NH₂
- THIQ-Leu-ACHPA-Leu-Phe-NH₂
- THIQ-Phe-ACHPA-Leu-Phe-NH₂
- THIQ-Val-ACHPA-Leu-Phe-NH₂
- THIQ-Cpa-ACHPA-Leu-Phe-NH₂
- 20 THIQ-Ala-ACHPA-Ala-Phe-NH₂
- THIQ-Val-ACHPA-Ala-Phe-NH₂
- THIQ-Val-ACHPA-Phe-Phe-NH₂
- THIQ-Val-ACHPA-Gly-Phe-NH₂
- THIQ-Val-ACHPA-Asn-Phe-NH₂
- 25 THIQ-Val-ACHPA-Gln-Phe-NH₂
- THIQ-Val-ACHPA-Asp-Phe-NH₂
- THIQ-Val-ACHPA-Glu-Phe-NH₂
- THIQ-Ala-ACHPA-Leu-Phe-OH
- THIQ-Ala-AHPPA-Leu-Phe-NH₂
- 30 THIQ-Val-AHPPA-Leu-Phe-NH₂
- THIQ-Ala-Sta-Leu-Phe-NH₂
- THIQ-Ile-ACHPA-Cha-Phe-NH₂
- THIQ-Val-ACHPA-Cha-Phe-NH₂
- THIQ-Ile-ACHPA-Leu-Phe-NH₂
- 35 THIQ-Val-Pheψ[CH₂NH]Leu-Leu-Phe-NH₂
- THIQ-Val-Chaψ[CH₂NH]Leu-Leu-Phe-NH₂
- THIQ-Val-Pheψ[CH₂NH]Leu-Gln-Phe-NH₂
- THIQ-Asn-Pheψ[CH₂NH]Leu-Gln-Ile-NH₂
- 3-(1,2,3,4-tetrahydro-7-nitroisoquinolyl)carbonyl-Val-ACHPA-Leu-Phe-NH₂ and
- 40 3-(7-amino-1,2,3,4-tetrahydroisoquinolyl)carbonyl-Val-ACHPA-Leu-Phe-NH₂

8. A pharmaceutical composition comprising a peptide as recited in any one of claims 1 to 7, or a therapeutically acceptable salt thereof, and a pharmaceutically acceptable carrier.

9. Use of a peptide as recited in any one of claims 1 to 7, or a therapeutically acceptable salt thereof for the preparation of a composition for treating HIV infections in a human.

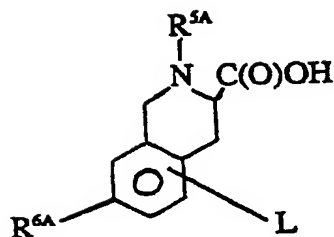
45 10. Use of a peptide as recited in any one of claims 1 to 7, or a therapeutically acceptable salt thereof for the preparation of a composition for protecting human cells against HIV pathogenesis.

11. Use of a peptide as recited in any one of claims 1 to 7, or a therapeutically acceptable salt thereof for the preparation of a composition for treating renin-associated hypertension in a mammal.

12. Use of a peptide as recited in any one of claims 1 to 7, or a therapeutically acceptable salt thereof 50 for the preparation of a composition for treating congestive heart failure in a mammal.

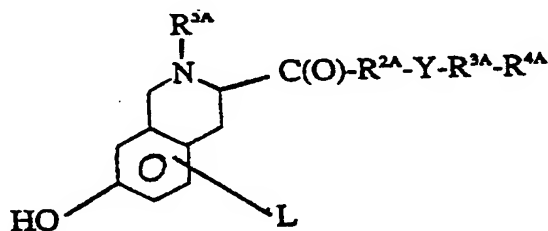
13. A process for preparing a peptide of formula 1 of claim 1 comprising:

(i) coupling the derived amino acid of formula 3

3

wherein R^{5A} is lower alkyl, or $QOC(O)-$ or $QC(O)-$ wherein Q is as defined in claim 1; R^{6A} is hydrogen, hydroxy, lower alkyl, lower alkoxy, amino, nitro or $OCH_2C(O)OR^{7A}$ wherein R^{7A} is lower alkyl, lower cycloalkyl, (lower cycloalkyl)methyl, phenyl, phenyl monosubstituted with lower alkyl, lower alkoxy or halo, benzyl or benzyl monosubstituted with lower alkyl, lower alkoxy or halo, and L is as defined in claim 1, with a fragment of formula $H-R^{2A}-Y-R^{3A}-R^{4A}$ wherein R^{2A} , R^{3A} and R^{4A} have the same meaning as defined in claim 1 for R^2 , R^3 and R^4 , respectively, except that side chain amino and carboxy groups, and a C-terminal carboxy group, if present, are replaced with corresponding protecting groups, and Y is as defined in claim 1; or

(ii) subjecting the intermediate of formula 4

4

wherein R^{2A} , Y, R^{3A} , R^{4A} and R^{5A} are as defined in this claim to O-alkylation with the alkylating agent of formula $XCH_2C(O)OR^{7A}$, wherein X and R^{7A} are as defined in this claim, in the presence of a strong base; followed, if required, by amino or carboxy deprotection, to give the corresponding peptide of formula 1; and, if desired, transforming the peptide into a therapeutically acceptable salt.



European Patent
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EUROPEAN SEARCH REPORT

Application Number

EP 90 11 0322

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.5)
X	EP-A-0 220 665 (MERCK) * Column 23, compound no. 17; claim 1 *	1,8,11-13	C 07 K 5/02
Y	---	2-6	
Y	TETRAHEDRON, vol. 44, no. 3, 1988, pages 661-673, Oxford, GB; T.K. SAWYER et al.: "Structure-conformation-activity relationships of renin inhibitory peptides having P1-P1'Xaaphi[CH2NH]Yaa substitutions: Molecular modeling and crystallography studies" * Page 663, table 1; page 666, table 2 *	2-6	
Y	PEPTIDES - STRUCTURE AND FUNCTION, PROCEEDINGS OF THE NINTH AMERICAN PEPTIDE SYMPOSIUM, ed. DEBER et al., 1985, pages 747-751, Pierce Chemical Co., Rockford, Illinois, US; J. BOGER et al.: "Renin inhibitors. Tripeptide and tetrapeptide competitive inhibitors containing a novel analog of statine" * Page 749, table I *	2-6	
			TECHNICAL FIELDS SEARCHED (Int. Cl.5)
			C 07 K
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 04-09-1990	Examiner KORSNER S.E.
CATEGORY OF CITED DOCUMENTS			
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons * : member of the same patent family, corresponding document	

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